

Prostate Short-Chain Dehydrogenase Reductase 1 (*PSDR1*): A New Member of the Short-Chain Steroid Dehydrogenase/Reductase Family Highly Expressed in Normal and Neoplastic Prostate Epithelium¹

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ABSTRACT

Genes regulated by androgenic hormones are of critical importance for the normal physiological function of the human prostate gland, and they contribute to the development and progression of prostate carcinoma. We used cDNA microarrays comprised of prostate-derived cDNAs to profile transcripts regulated by androgens in prostate cancer cells. This study identified a novel gene that we have designated prostate short-chain dehydrogenase/reductase 1 (*PSDR1*), that exhibits increased expression on exposure to androgens in the LNCaP prostate cancer cell line. Northern analysis demonstrated that *PSDR1* is highly expressed in the prostate gland relative to other normal human tissues. The *PSDR1* cDNA and putative protein exhibit homology to the family of short-chain dehydrogenase/reductase enzymes and thus identify a new member of this family. Cloning and analysis of the putative *PSDR1* promoter region identified a potential androgen-response element. We used a radiation-hybrid panel to map the *PSDR1* gene to chromosome 14q23-24.3. *In situ* hybridization localizes *PSDR1* expression to normal and neoplastic prostate epithelium. These results identify a new gene involved in the androgen receptor-regulated gene network of the human prostate that may play a role in the pathogenesis of prostate carcinoma.

INTRODUCTION

Prostate adenocarcinoma is responsible for more than 39,000 deaths annually in the United States (1). Circulating androgens and the intracellular AR³ are critical mediators of prostate cancer growth and the progression to lethal disease. Landmark discoveries by Huggins and Hodges in 1941 (2) demonstrated that most prostate cancers are initially androgen-dependent, a finding that initiated the era of effective endocrine-based therapy for this malignancy. To date, surgical or chemical castration remains the mainstay of therapy for advanced prostate cancer. A reduction in serum testosterone leads to marked tumor regression through a mechanism of programmed cell death (3). Although responses to this therapy may last for years, the approach is rarely curative because surviving cancer cells lose their dependency on exogenous testosterone over time and are capable of proliferating in the absence of detectable serum androgens.

In addition to a role in driving cellular proliferation, an intact androgen signaling system may also be associated with tumor sup-

pression (4). This dual role of androgens would not be unexpected, because androgens are responsible for differentiation of the prostate epithelium and for the regulation of specific epithelial cell functions such as the expression of PSA (5, 6). Several androgen-regulated genes have been demonstrated to be associated with a proliferative shut-off function in LNCaP cells and for the regulation of the cell cycle (7, 8). At the time of invasion or metastasis, mutations in the AR may occur (9), suggesting that a normal AR is protective from progression. Finally, *in vitro* studies indicate that there may be a survival advantage in maintaining an androgen-responsive cohort of prostate tumor cells (10). This concept has been extended to clinical medicine in which several trials suggest a benefit for an approach using intermittent rather than continuous androgen suppression in patient cohorts with hormone-responsive disease (11, 12).

The pivotal role of androgens in the biology and treatment of prostate cancer has led to intensive investigations designed to identify the molecular mediators of androgen action (13, 14). Among the genes shown to be regulated by androgens in prostate cells are several that encode enzymes belonging to the two major lipogenic pathways: fatty acid synthesis and cholesterol synthesis (15, 16). The regulation of cholesterol metabolism by androgens is especially intriguing because cholesterol is an essential precursor for the biosynthesis of androgens (17). Other molecules involved in androgen metabolism and androgen action are themselves androgen regulated. For example, 17- β -HSD, an enzyme that converts androstenedione to testosterone, is androgen regulated (18), as is the expression of the AR itself (7, 19). It appears that multiple autoregulatory levels of androgen action may be operative in androgen-responsive tissues.

Our objective in this study was to identify genes that exhibit transcriptional regulation by androgens in human prostate cells. We hypothesized that such genes could be direct mediators of androgen action and that the characterization of these genes and their cognate proteins would provide insights into the mechanisms of androgen-dependent and androgen-independent cellular growth. We used cDNA microarrays comprised of cDNAs derived from human prostate tissues to quantitate transcripts expressed in the androgen-sensitive LNCaP prostate tumor cell line under conditions of androgen starvation or androgen stimulation. Here we report the cDNA cloning, chromosomal mapping, genomic structure, and expression profile of a novel gene, *PSDR1*, that exhibits homology to the family of SDR enzymes. *PSDR1* is the first SDR shown to be predominantly expressed in normal and neoplastic prostate tissue. We hypothesize that *PSDR1* may play a role in steroid hormone metabolism in prostate cells and thus may be an ideal target for modulating hormone-mediated prostate cancer growth.

MATERIALS AND METHODS

Microarray Fabrication. A nonredundant set of 1500 prostate-derived cDNA clones were identified from the Prostate Expression DataBase (PEDB), a public sequence repository of EST data derived from human prostate cDNA

Received 6/7/00; accepted 12/13/00.

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¹ Supported in part by the CaPCURE Foundation, by Grant CA75173-01A1 from the National Cancer Institute (NIH) (to P. S. N.), and by gifts from the Seattle Foundation and the Lazar Foundation.

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³ The abbreviations used are: AR, androgen receptor; PSA, prostate-specific antigen; ARE, androgen response element; DHT, dihydrotestosterone; EST, expressed sequence tag; dbEST, database of ESTs; CS-FCS, charcoal-stripped FCS; PRE, progesterone responsive element; HSD, hydroxysteroid dehydrogenase; RACE, rapid amplification of cDNA ends; SDR, short-chain dehydrogenase reductase; *PSDR1*, prostate SDR 1; hk, human glandular kallikrein.

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libraries (20). Individual clone inserts were amplified by the PCR using 2 μ l of bacterial transformant culture as template with primers BL_m13F (5'-GTAAACGACGGCCAGTGAATTG-3') and BL_m13R (5'-ACACAGGAAACAGCTATGACCATG-3') as described previously (21). PCR products were purified through Sephadryl S500 (Pharmacia), mixed 1:1 with DMSO (Amersham), and spotted in duplicate onto coated type IV glass microscope slides (Amersham) using a Molecular Dynamics GenII robotic spotting tool. After spotting, the glass slides were air-dried and UV-cross-linked with 500 mJ of energy and then baked at 95°C for 30 min.

Probe Construction and Microarray Hybridization. Total RNA was isolated from LNCaP cells after 72 hrs of androgen depletion or supplementation using TRIzol (Life Technologies) according to the manufacturer's directions. Fluorescence-labeled probes were made from 30 μ g total RNA in a reaction volume of 20 μ l containing 1 μ l of anchored oligo-dT primer (Amersham), 0.05 mM Cy3-dCTP or Cy5-dCTP (Amersham); 0.05 mM dCTP; 0.1 mM each dGTP, dATP, dTTP; and 200 units of Superscript II reverse transcriptase (Life Technologies). Reactants were incubated at 42°C for 120 min followed by heating to 94°C for 3 min. Unlabeled RNA was hydrolyzed by the addition of 1 μ l of 5 N NaOH and heating to 37°C for 10 min. One μ l of 5 M HCl and 5 μ l of 1 M Tris-HCl (pH 7.5) were added to neutralize the base. Unincorporated nucleotides and salts were removed by chromatography (Qiagen), and the cDNA was eluted in 30 μ l of dH₂O. One μ g of dA/dT 12-18 (Pharmacia) and 1 μ g of human CotI DNA (Life Technologies) were added to the probe, heat denatured at 94°C for 5 min, combined with an equal volume of 2 \times microarray hybridization solution (Amersham), and prehybridized at 50°C for 1 h. The mixture was then placed onto a microarray slide with a coverslip and hybridized in a humid chamber at 52°C for 16 hours. The slides were washed once with 1 \times SSC, 0.2% SDS at room temperature for 5 min, then twice with 0.1 \times SSC, 0.2% SDS at room temperature for 10 min. After washing, the slides were rinsed in distilled water to remove trace salts and dried.

Image Acquisition and Data Analyses. Fluorescence intensities of the immobilized targets were measured using a laser confocal microscope (Molecular Dynamics). Intensity data were integrated at a pixel resolution of 10 μ m using ~20 pixels per spot, and recorded at 16 bits. Quantitative data were obtained with the SpotFinder V 2.4 program written at the University of Washington. Local background hybridization signals were subtracted prior to comparing spot intensities and determining expression ratios. For each experiment, each cDNA was represented twice on each slide, and the experiments were performed in duplicate producing four data points per cDNA clone per hybridization probe. Intensity ratios for each cDNA clone, hybridized with probes derived from androgen-stimulated LNCaP and androgen-starved LNCaP, were calculated (stimulated intensity/starved intensity). Gene-expression levels were considered significantly different between the two conditions if all four of the replicate spots for a given cDNA demonstrated a ratio >2 or <0.5, and the signal intensity was greater than 2 SD above the image background.

Cell Culture and General Methods. DNA manipulations including transformation, plasmid preparation, gel electrophoresis, and probe labeling, were performed according to standard procedures (22). The LNCaP prostate carcinoma cell line was cultured in RPMI 1640 supplemented with 10% FCS (Life Technologies, Rockville, MD). Cells were transferred into RPMI 1640 with 10% CS-FCS (Life Technologies) 24 h before androgen-regulation experiments. This medium was replaced with fresh CS-FCS media or CS-FCS supplemented with 1 nM synthetic androgen R1881 (NEN Life Science Products Inc.). Cells were harvested for RNA isolation at 0-h and 72-h time points.

Northern Analysis. Ten μ g of total RNA were fractionated on 1.2% agarose denaturing gels and transferred to nylon membranes by a capillary method (22). The human multiple tissue and master blots were obtained from Clontech. Blots were hybridized with DNA probes labeled with [α -³²P]dCTP by random priming using the Rediprime II random primer labeling system (Amersham) according to the manufacturer's protocol. Filters were imaged and quantitated by using a phosphor-capture screen and Image-quant software (Molecular Dynamics).

cDNA Library Screening and RACE. We screened 1,200,000 phage plaques from a human prostate 5'-stretch cDNA library (Clontech) with the 6A4 cDNA probe representing the 3' end of the PSDRI cDNA. Two separate rounds of library screening identified 16 partial-length cDNA clones. Searches of dbEST identified seven IMAGE cDNA clones (IMAGE CloneID: 360400, 109237, 1130518, 1401718, 1337270, 1723130, 1703429) that contained se-

quences homologous to PSDRI. All of the clones were sequenced and assembled using the Sequencer software (Gene Codes, Corp.). To clone the 5' end of the cDNA, 5'-RACE was performed on human prostate Marathon-Ready cDNA (Clontech) using primers 6A3_RC3 (5'-GGACAGCATTTTCCTGATTGGGGC-3') and 6A4_RC4 (5'-CAGAAGGAGGAGCAACAGCGGGAAC-3'). The RACE products were subcloned into PCR2.1-TOPO (Invitrogen) and sequenced.

Phage plaques (1,200,000) from a human prostate 5'-STRETCH cDNA library (Clontech) were screened with PSDRI ³²P-cDNA probes according to the manufacturer's instructions. Eleven additional cDNA clones were isolated, subcloned and sequenced. RACE reactions were performed using the human prostate Marathon-ready cDNA cloning kit (Clontech) following the manufacturer's instructions. Templates for RACE reactions were prostate Marathon-ready cDNA (Clontech) and androgen-stimulated LNCaP cDNA prepared using Marathon cDNA amplification kit (Clontech). Nested 5'-RACE reactions were performed according to the manufacturer's instructions; first with primers 6A4RC4, 5'-CAGAAGGAGGAGCAACAGCGGGAAC-3' and AP1 (Clontech) and then a nested RACE reaction with primers 6A4RC3, 5'-GGACAGCATTTTCCTGATTGGGGC-3' and AP2 (Clontech). The RACE products were subcloned into PCR2.1-TOPO vectors with the TOPO TA cloning kit (Invitrogen) and sequenced.

Chromosomal Localization of PSDRI by Radiation Hybrid Panel Mapping. The G3 Gene bridge radiation hybrid panel (Research Genetics, Huntsville, AL) was used to map the chromosomal localization of PSDRI with primers 6A4F (5'-GGGGCATTTTCCTTACATTGTCTTG-3') and 6A4R (5'-CACTCCAAACAAGTGATGGGAACAC-3'). After 35 cycles of amplification, the reaction products were separated on a 1.2% agarose gel, and the resulting product pattern was analyzed through the Stanford genome center web server⁴ to determine the probable chromosomal location.

In Situ Hybridization. For mRNA *in situ* hybridization, recombinant plasmid pCRII-TOPO (Invitrogen), containing a 400-bp PSDRI fragment was linearized to generate sense and antisense digoxigenin-labeled RNA probes. *In situ* hybridization was performed according to the manufacturer's protocol on the Ventana GenII automated instrument (Ventana Medical Systems, Tucson, AZ). Tissue sections (5 μ m) were mounted onto Chroma plus slides (VWR Scientific), deparaffinized in a 65°C oven for 2 h followed by three 5-min soaks in xylene and rehydrated through graded alcohol with a final rinse in 2 \times SSC. Before hybridization, sections were digested with proteinase I cocktail for 12 min at 37°C, then 10 ng of either sense or antisense probe in the hybridization buffer was applied. Programmed recipe files consisting of buffer rinses, protease digestion, hybridization, detection, and counterstains were optimized for the PSDRI probe. Digoxigenin-labeled RNA probe was added manually. Antidigoxigenin was used as the primary antibody. The probe was denatured at 65°C, and hybridization was carried out at 42°C for 360 min. Washes were performed at 37°C with 2 \times , 1 \times , and 0.1 \times SSC. The system uses a cocktail of antirabbit and antimouse secondary IgG-biotinylated antibody with an indirect biotin-avidin diaminobenzidine detection system. The sections were counterstained with hematoxylin.

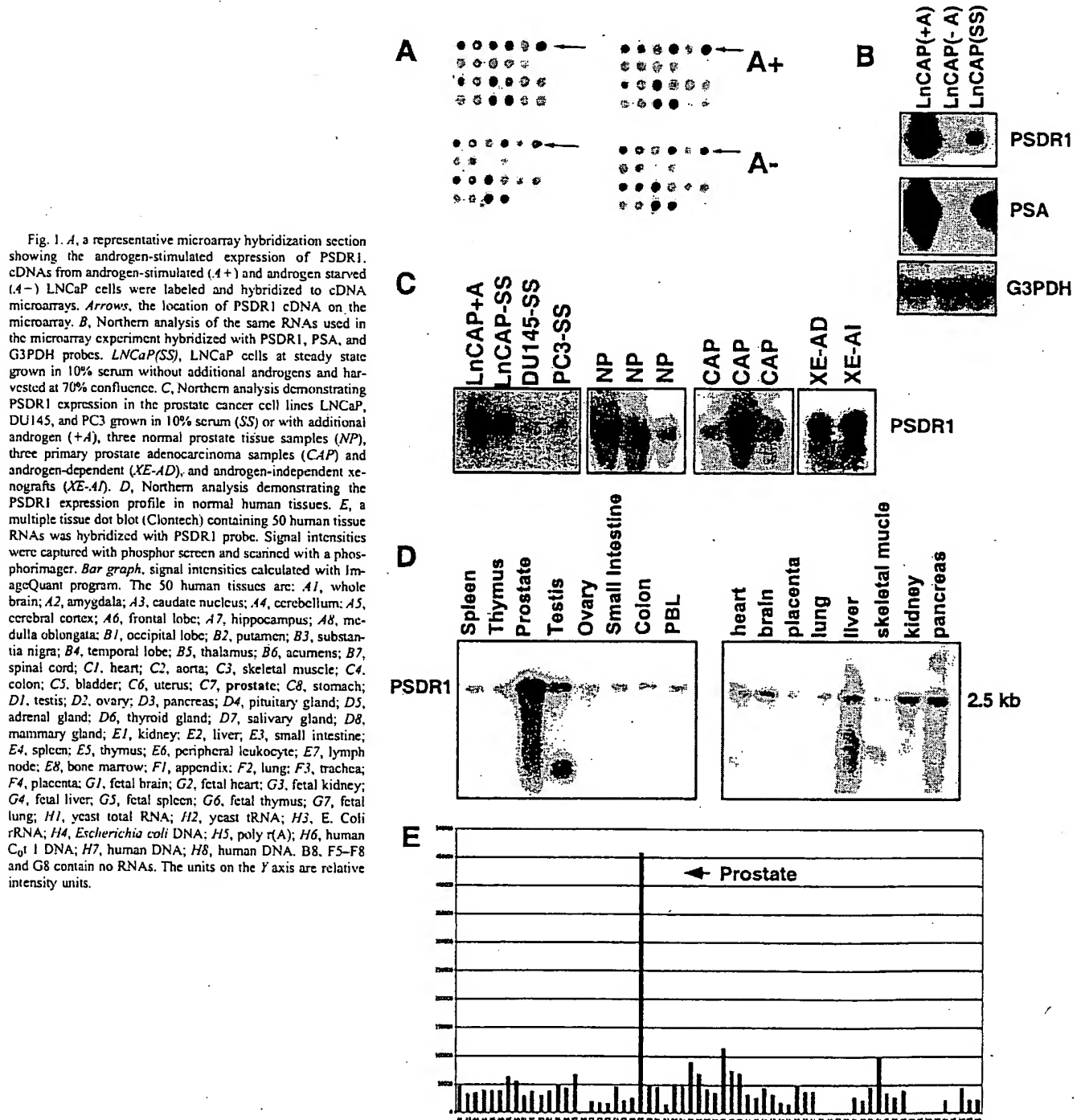
RESULTS

Identification of a Novel Androgen-regulated cDNA, PSDRI, by Microarray Expression Analysis. Microarrays comprised of cDNA clones derived from prostate tissues were hybridized with total cDNA probes synthesized from androgen-stimulated and androgen-starved LNCaP prostate cancer cells. Four independent data points for each arrayed cDNA were generated. The hybridization ratios for 20 distinct cDNAs were consistently increased by >2-fold in androgen-stimulated relative to androgen-starved cells. We did not observe any cDNAs with consistent hybridization ratios <0.5, a ratio that would indicate down-regulated expression. The genes induced by androgens included *hK2* (23), *hK3*, also known as *PSA* (24), *NKX3.1* (25), *prostate/PRSS17* (26), *TMPS2* (13), *PART-1* (27), several genes involved in lipid metabolism, and several anonymous ESTs. The expression level of

⁴ Internet address: <http://shgc.stanford.edu>.

the cDNA clone corresponding to one of these ESTs, 6A4, increased 3-fold in androgen-stimulated LNCaP cells relative to androgen-deprived cells as assayed by microarray hybridization (Fig. 1A). Sequence comparisons against the GenBank and dbEST databases revealed homology only to uncharacterized partial-length ESTs (e.g., AA657851, IMAGE ID:1207405). Full-length cloning of the corresponding cDNA and subsequent nucleotide and amino acid sequence comparisons revealed significant homology to conserved motifs of the SDR family of proteins. We have named this gene *PSDR1* for Prostate Short-chain Dehydrogenase/Reductase 1.

Cloning of the Full-Length *PSDR1* cDNA. We screened a human prostate cDNA library with the 6A4 cDNA probe representing the 3' end of the *PSDR1* cDNA and identified 16 partial-length cDNA clones. Searches of the dbESTs initially identified seven IMAGE cDNA clones (IMAGE Clone ID: 360400, 109237, 1130518, 1401718, 1337270, 1723130, 1703429) that contained sequence homologous to *PSDR1*. To clone the 5' end of the cDNA, 5'-RACE was performed on cDNA from normal human prostate (Clontech) and the LNCaP cell line. All of the clones were sequenced and subsequently assembled using the Sequencer software (Gene Codes, Corp.). A total of 2539 bp were obtained, which corresponds to the 2.5-kb band



which is in agreement with the size of its transcript as determined by Northern hybridization (Fig. 1D). The cDNA sequence was submitted to GenBank under the accession no. AF167438.

The *PSDR1* cDNA encodes a putative protein of 318 amino acids. The start codon, GAGATGG matches in a strong context to the Kozak translation initiation consensus sequences (RNNATGG, where R is a purine; Ref. 28). Two potential polyadenylation signals were identified at nucleotide positions 2439 and 2481. IMAGE clone 1703429 has a poly(A) stretch that uses the AATAAA polyadenylation signal at 2419, and our original cDNA clone 6A4 uses the AATAAA signals at 2481. However, we were not able to find a polyadenylation site that would produce the 900-bp band seen in testis tissue. PCR primers flanking the start and stop codons were designed, and an expected size band encompassing the entire coding region was amplified from human prostate Marathon-Ready cDNA (Clontech; data not shown).

Comparisons of the assembled cDNA sequences indicated several polymorphic sites. Five distinct single nucleotide polymorphisms were recognized between the three independent prostate tissue sources used for *PSDR1* cloning. Three occur in the coding region of the *PSDR1* sequence; nucleotide 379, ggc to ggg; nucleotide 916, gtg to gtc; and nucleotide 921, gtc to gcc. The first two are conserved changes, whereas the latter results in a valine to alanine amino acid substitution. Alignments of sequences in dbEST with homology to *PSDR1* identified more than 20 distinct nucleotide differences in tissue sources presumably derived from different individuals.

Prostate-localized and Androgen-regulated Expression of *PSDR1*. The androgen-regulated expression of *PSDR1* was confirmed by Northern analysis using the same LNCaP RNA that was used for microarray analysis. PhosphorImage quantitation of the Northern analysis demonstrated a 3-fold induction of *PSDR1* expression after 72 h of androgen exposure relative to 72 h of androgen starvation (Fig. 1B). PSA expression increased 25-fold, and the expression of the G3PDH loading control did not change significantly. Interestingly, Northern analysis with androgen-independent prostate cancer cell lines DU145 and PC3 demonstrated *PSDR1* expression in both cell types (Fig. 1C) indicating a mechanism of *PSDR1* transcription in these cells that is independent of androgen requirements.

The distribution of *PSDR1* transcripts in normal human tissues and prostate carcinoma was determined by Northern analysis and mRNA dot blot. Of 16 adult tissues examined by Northern, a *PSDR1* message of 2.5 kb was predominantly expressed in prostate (Fig. 1D). In testis, the *PSDR1* probe hybridized to an additional band at about 900 bp (Fig. 1D), which could indicate cross-hybridization, alternate splicing, or alternate usage of polyadenylation signals. The *PSDR1* expression profile was confirmed using an RNA Master dot blot (Clontech) comprised of RNA from 50 different tissues. *PSDR1* expression was detected predominantly in prostate with a very low relative level of expression in spleen, thymus, testis, ovary, small intestine, colon, peripheral blood leukocyte, and kidney, adrenal gland, and fetal liver (Fig. 1E). *PSDR1* expression was at least 4-fold higher in prostate relative to any other human tissue examined. *PSDR1* expression was detected in all of the normal and neoplastic prostate tissue samples examined. These included three normal whole prostate tissues; three primary prostate adenocarcinomas, androgen-dependent and androgen-independent prostate cancer xenografts; and three prostate cancer cell lines (Fig. 1C).

***PSDR1* Shares Homology with Members of the SDR Family.** We used the nucleotide and translated the 318-amino acid *PSDR1* sequence to search the National Center for Biotechnology Information sequence databases by using BLAST and BEAUTY algorithms (29). Partial homology was seen with several oxidoreductases from bacteria and plant sources. To see whether the homology was significant, we searched the protein sequence of *PSDR1* against the BLOCKS data-

base (30).⁵ The *PSDR1* protein has three blocks that all match to the SDR family protein signature BLOCK (BL00061; Ref. 31) with a significant combined E-value of 2.6e-06. The SDR family are NAD- or NADP-dependent oxidoreductases (31), which include enzymes involved in steroid metabolism such as estradiol 17- β -dehydrogenase (also called 17- β -hydroxysteroid dehydrogenase; EC 1.1.1.62), 15-hydroxyprostaglandin dehydrogenase (NAD+) (EC 1.1.1.141) from human and 11 β -HSD (EC 1.1.1.146; 11-DH; Ref. 31). A multiple sequence alignment of the *PSDR1* protein with different members of the human HSD family and a prokaryotic 20- β -HSD (*Streptomyces* 3 α /20 β -HSD) is shown in Fig. 2.

Only two motifs are highly conserved in the SDR family. The first is a common GlyXXXGlyXGly pattern, in which the coenzyme NAD(H) or NADP(H) binds at the NH₂ terminus of the SDR enzyme (31). The second motif is a segment, TyrXXXLYs, believed to be involved in the catalytic activity of the enzyme (32). The *PSDR1* protein contains these two signatures (as shown by asterisk in Fig. 2). Sequence alignments reveal that proteins in the SDR family exhibit residue identities of only about 15–30%, probably because of their early divergence and remote origin (31). *PSDR1* shows ~25% amino acid identity with other members of the SDR family.

Searches against prosite patterns database⁶ revealed that *PSDR1* contains two Asn-glycosylation sites at amino acid (aa) position 174 and 198. These two sites are also conserved among SDR family proteins (Fig. 2). In addition, two protein kinase C (PKC) phosphorylation sites (aa 57 and 106), a casein kinase II phosphorylation site (aa 57), and a 7 N-myristoylation site are identified in the protein.

***PSDR1* Genomic Organization and Promoter Sequence Analysis.** BLAST searches with the full-length *PSDR1* cDNA identified homology with nucleotide sequence derived from a recently deposited unannotated 197-kb chromosome 14 BAC clone, R-1012A1, sequenced by the National Sequencing Center-Genoscope in France (GenBank accession no. AL049779). Alignment with the *PSDR1* sequence demonstrated that this BAC contains the entire *PSDR1* cDNA and allowed for the determination of the *PSDR1* genomic structure. The *PSDR1* gene comprises 7 exons and 6 introns. The sizes of exons and introns and the exon/intron junctional sequences are listed in Fig. 3B. All of the intron/exon junctions conform to the 5'-gt...3'-ag consensus (33) except intron 2. Intron 2 has a 5'-gc...3'-ag splicing signal, a structure that has been identified in other genes (34).

We examined the 5' genomic sequences for potential transcriptional start sites using a neural network promoter prediction program⁷ (35) and for potential transcriptional factor binding site using the TESS (Transcription Element Search Software) program⁸ (36). We identified a strong promoter sequence with a score of 0.87 (a score of 0.85 has a 0.1–0.4% false positive prediction rate). The predicted transcription start site is 167 bp 5' of the ATG start codon. A TATA box (TATAAT) is found 30 bp 3' of the putative transcriptional initiation site (Fig. 3A). A sequence that has 86.7% homology (13 of 15 nucleotides) to the consensus ARE, 5'-GGA/TACAnnnTGTTCT-3', (37) was identified (Fig. 3A). Two sequences that have 86.7% (13 of 15 nucleotides) homology to the consensus sequence of PREs (38) were also identified (Fig. 3A). An interleukin-6 response element binding protein site, TTCCAGAA, (39) was identified 281 bp 5' of the transcription initiation site.

Chromosomal Localization of *PSDR1*. The medium-resolution Stanford G3 radiation hybrid panel was used to determine the chro-

⁵ Internet address: <http://www.blocks.fhcrc.org>.

⁶ Internet address: <http://www.isrec.isb-sib.ch/software/PSTSCAN> form.html.

⁷ Internet address: <http://www.hgc.lbl.gov/projects/promoter.html>.

⁸ Internet address: <http://www.cbil.upenn.edu/less/index.html>.

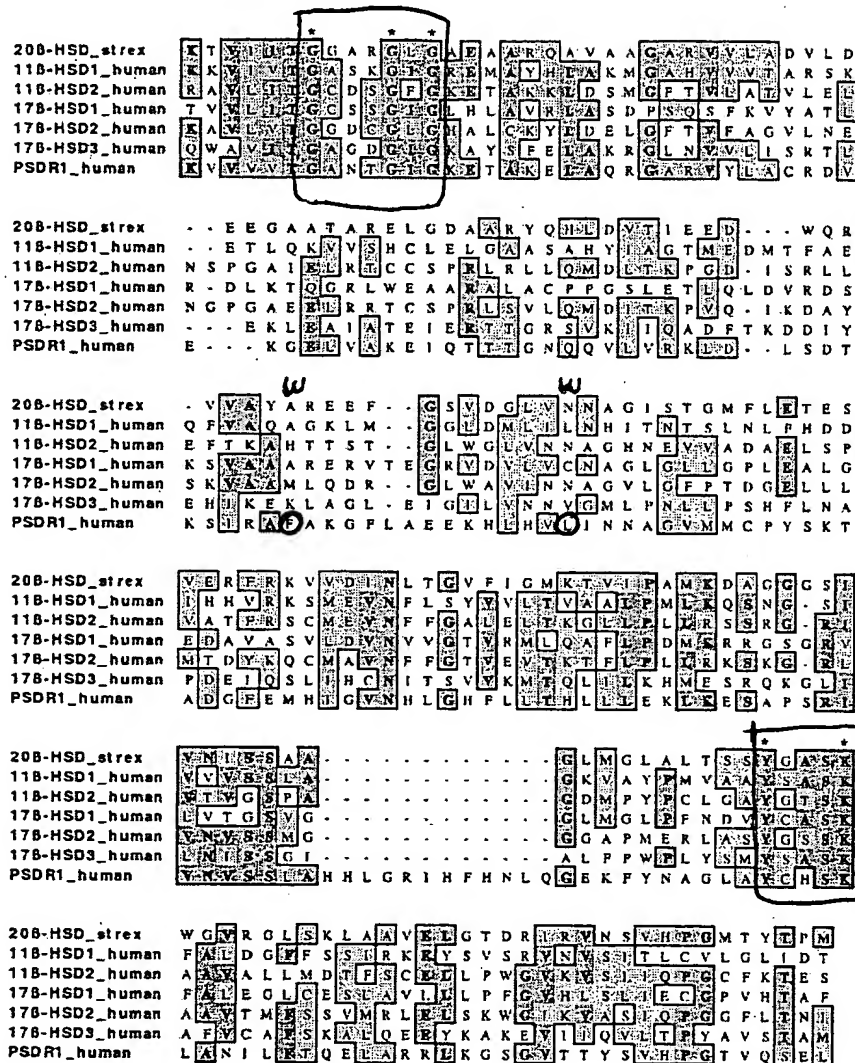


Fig. 2. Multiple sequence alignments of PSDR1 with different members of the human HSDs from the SDR family. A prokaryotic 20 β -HSD (*Streptomyces 3 α /20 β -HSD*) was included at the top (20- β HSD_strex). The alignment was performed with the clustalW algorithm (56) using MacVector 6.0 software (Oxford Molecular). BLOSUM series matrix was used with an open-gap penalty score of 10 and extend-gap penalty score of 0.05. Boxed and dark-shaded; identical residues; boxed and light-shaded, similar residues. *, two conserved segments of the SDR family. GlyXXXGlyXGly and TyrXXXLys. The GenBank accession numbers for members aligned here are: 20- β HSD_strex, *Streptomyces 3 α /20 β -HSD*, P19992; 11- β HSD1_human, P28845; 11- β HSD2_human, U14631; 17- β HSD1_human, P14061; 17- β HSD2_human, L11708; 17- β HSD3_human, P37058. Only the regions containing the conserved motifs are shown here.

mosomal localization of *PSDR1* using gene-specific PCR primers 6A4F and 6A4R. Analysis of the typing results on the Stanford Human Genome Center Radiation Hybrid Panel server⁹ indicated that *PSDR1* is located closest to Stanford Human Genome Center Radiation Hybrid Panel-2558 between two cytogenetically mapped markers D4S63 (mapped to 14q23) and D4S258 (mapped to 14q24.3).⁹ Therefore, *PSDR1* is mapped to 14q23-24.3, consistent with the BAC clone mapping data localizing *BAC R-1012A1* to chromosome 14q.

***PSDR1* Expression in Normal and Neoplastic Prostate Epithelium.** Normal prostate contains two major epithelial cell populations, the luminal secretory cells and the basal cells. *In situ* hybridizations were performed on sections of normal prostate by using an antisense RNA probe specific for *PSDR1* to localize its expression. *PSDR1* was expressed in both normal basal and luminal cell populations. (Fig. 4, A and C). Little to no staining was seen in fibromuscular stromal cells, endothelial cells, or infiltrating lymphocytes. Hybridization with sense *PSDR1* RNA probes showed no background staining (Fig. 4, B and D). *In situ* hybridizations with *PSDR1* antisense and sense probes were also performed on sections of primary prostate adenocarcinoma obtained from radical prostatectomy specimens. Adenocarcinoma cells were uniformly positive for *PSDR1* expression (Fig. 4E). Hy-

bridization with sense *PSDR1* RNA probes showed no background staining (Fig. 4F).

DISCUSSION

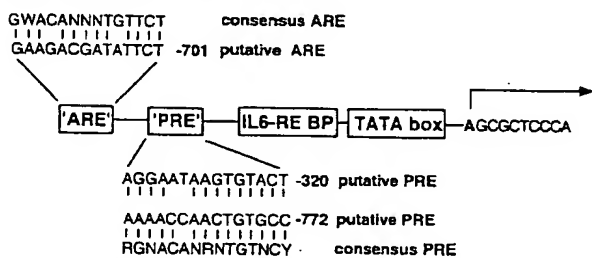
In a search for genes regulated by androgens in the human prostate, we have identified a new member of the SDR superfamily. SDRs encompass a large group of functionally diverse proteins in pro- and eukaryotes (31). Enzymes in this family typically exhibit residue identities of only 15–30%, indicating early gene duplication events and subsequent extensive divergence (31). Regions of high conservation are restricted to specific segments, which indicate a possible common fold, active site, reaction mechanism, and coenzyme and substrate binding regions (40). Of relevance for the study of androgen-mediated effects in prostate carcinoma is the classification of several key enzymes involved in steroid biosynthesis, HSDs, within the SDR family. This group of HSDs includes 17- β -HSD types 1–4 and 6 (41), 15-hydroxyprostaglandin dehydrogenase, and 11- β -HSD (31). 17- β -HSD 3 converts androstenedione to testosterone (42); 17- β -HSD 6 converts 5-androstane-3 α , 17- β -diol (3- α -diol) to androsterone (43). In prostate cancer cells, 17- β -HSD type 2 exclusively converts 5 α -DHT and testosterone into the less potent 17-keto compounds 5 α -androstanedione and 4-androstenedione, respectively (44). This suggests

⁹ The Genome Database: <http://www.gdb.org/>.

that 17HSD type 2 plays a part in the androgen metabolic pathway, resulting in the inactivation of testosterone and 5 α -DHT locally in the prostate. Enzyme expression in the prostate could, therefore, protect cells from excessive androgen action. Because *PSDR1* shows significant homology to HSD members of the SDR family, we hypothesize that *PSDR1* is also involved in prostate cellular steroid metabolism that may include biosynthesis and/or degradation. The pharmacological modulation of *PSDR1* activity could thus influence prostate cellular growth and provide a new target for prostate cancer therapy.

Numerous studies support associations between molecular variations involving genes of the androgen metabolic pathway and the development and progression of prostate cancer (45). In addition to environmental influences, racial and international variation in prostate cancer incidence suggests that inheritable genetic factors such as those that influence androgen biosynthesis, activation, transport, and metabolism are operative (45). In addition to polymorphic variation in the AR itself (46), specific polymorphisms in the 5 α -reductase type 2 (*SRD5A2*) gene, the enzyme converting testosterone to the more bioactive DHT, result in increased enzyme activity and confer up to a 7-fold increased risk for the development of prostate cancer in African-American men (47). Allelic variants in the 3- β -HSD type II gene, encoding one of two enzymes that initiates the inactivation of DHT, have been identified and are currently under assessment for a role in racial/ethnic differences in prostate carcinogenesis (48). Polymorphisms in *PSDR1* could influence enzyme activity and consequently result in variations in steroid metabolism between individuals. Our preliminary analysis of the *PSDR1* sequence from three different prostate tissue sources identified five distinct single nucleotide polymorphisms. Three occur in the coding region of the *PSDR1* sequence, one of which results in a valine to alanine amino acid substitution. An alignment of sequences in dbEST with homology to *PSDR1* identified more than 20 distinct nucleotide differences in tissue sources presumably derived from different individuals. Although some of these differences may represent sequencing artifacts, these findings warrant a more directed study of *PSDR1* variation in different ethnic populations and in samples of prostate carcinoma.

A. Schematic of the PSDR1 promoter region.



B. Summary of the genomic structure of PSDR1.

Exon	Acceptor	Donor	Exon size	Intron size (bp)
1	—	CAGgtctgtgcaaatgtatgcc	>114	2577
2	ctctcctctctgtctgcagGAA	GAGgcaagltcaocctcttcaa	120	340
3	ttcatatgttgctgacagGAG	GAGgtaagltgtagaactagaga	159	1193
4	tcgtctgttccctgcagagGAA	TGGgtaagaaatctggcctatc	104	718
5	attctagttattctcaacagGTC	AAGgtggcctagagaaatga	211	5007
6	atgccacccccaaccagGCT	CAGgtatgaatgttatctt	191	6591
7	ccttctcttaccctcagTGA	—	1621	—

Fig. 3. Genomic structure of PSDR1. A, schematic drawing showing the putative sequence motifs of the PSDR1 promoter. Arrow, the predicted transcription initiation site A, the +1 position. TATAAT TATA-box at -30, putative ARE and PRE sequences, and an interleukin-6 response element binding protein IL-6 RE-BP site TTCCCAGAA at -281 are also shown. IUPUC-IUB codes for nucleotides were used: R, purine; Y, pyrimidine; W, A or T; B, not A (C, G, or T); K, G or T; M, A or C; N, any nucleotides. B, PSDR1 exon and intron acceptor and donor splice sites with corresponding segment sizes in nucleotide base pairs.

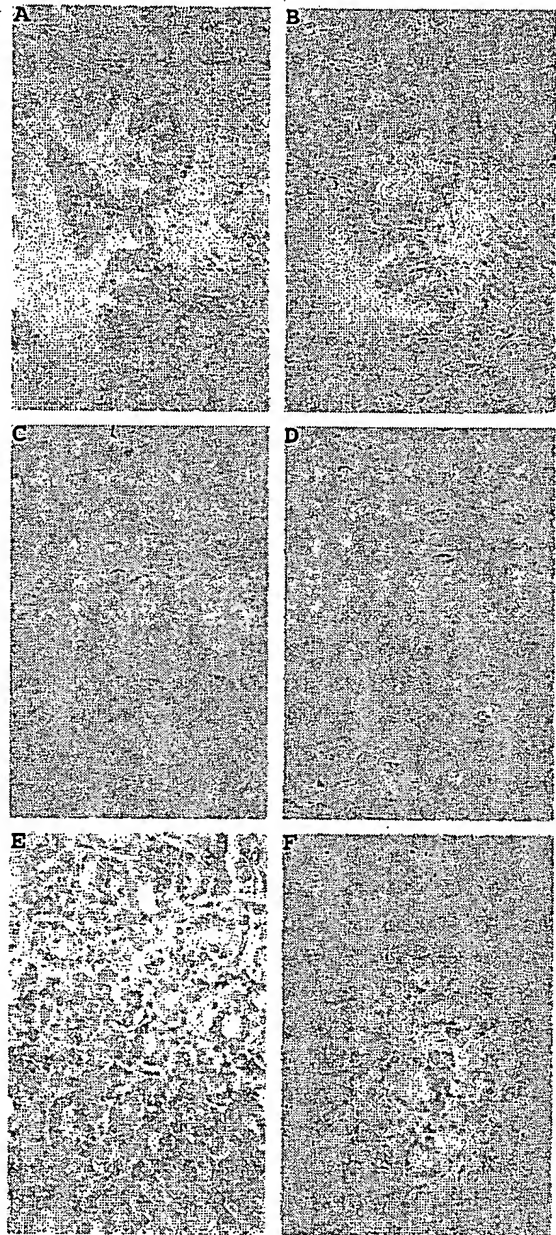


Fig. 4. Representative sections of *in situ* hybridization with PSDR1 probes in normal and malignant prostate tissues. Low (A) and high (C) power magnification of normal prostate tissue hybridized with PSDR1 antisense probe, showing expression in both basal and luminal cells, but not in stroma cells. Low (B) and high (D) power magnification of negative control hybridization with PSDR1 sense probes, showing no background staining in normal prostate tissue. E, primary prostate carcinoma tissue hybridized with PSDR1 antisense probe showing PSDR1 expression in prostate carcinoma cells. F, negative control hybridization with PSDR1 sense probes, showing no background tissue hybridization.

The expression of *PSDR1* is induced by synthetic androgens in LNCaP cells. The mechanism of androgen-mediated regulation of *PSDR1* expression is unknown and could involve either direct AR binding to *PSDR1* promoter regions or indirect activation through the modulation of intermediary transcription factors or via posttranscriptional mechanisms. Androgens have been shown to regulate expression of other oxidoreductases. The mouse alcohol dehydrogenase *ADH1*, which belongs to the long-chain dehydrogenase family, is induced 10–12 fold by androgens in mouse kidney cells (49). The induction of mouse *ADH1* gene by androgens seems to be AR de-

pendent because the *ADH1* gene in *Tfm* mice lacking functional AR was not responsive to androgens (49). 17- β -HSD1, which belongs to the SDR family, is stimulated by androgen through AR-mediated mechanism (50). We have identified a putative ARE and two putative PRE sites that demonstrate a high degree of homology to the respective consensus hormone receptor binding sites. Whether these responsive elements are functional awaits further investigation. In addition, *PSDR1* transcription is not entirely mediated by androgens as demonstrated by a low level of detectable *PSDR1* message in androgen-starved LNCaP cells and in the androgen-independent PC3 and DU145 prostate cancer cell lines. Studies of the *PSDR1* protein may identify additional mechanisms of functional regulation.

The localized expression of *PSDR1* in prostate epithelium is interesting because, to our knowledge, *PSDR1* is the first member of the human SDR family that is expressed predominantly in the prostate gland. Other members of SDR family exhibit tissue-restricted patterns of expression. For example, 17- β -HSD1 is predominantly expressed in placenta and ovary (51, 52); 17- β -HSD2 is expressed in placenta and liver (53); 17- β -HSD3 is expressed in testis (42); 17- β -HSD4 is primarily expressed in liver and kidney (54); 17- β -HSD5 is expressed most abundantly in liver and testis (41, 55); and 17- β -HSD6 is expressed equally in liver and prostate (43).

The identification of genes with selective expression in specific organs or cell types provides an entry point for understanding biological processes that occur uniquely within a particular tissue. Genes and their cognate proteins whose expression is specific for the prostate have greatly aided the diagnosis and treatment of prostate carcinoma. The significance of the prostate predominant expression pattern of *PSDR1* remains to be determined. If the tissue expression profile of the *PSDR1* protein corresponds to the transcript expression profile, then *PSDR1* may represent an additional target for prostate cancer diagnostic and therapeutic interventions. Cellular or humoral immunotherapy could be designed to exploit the tissue expression differential. Our hypothesis is that *PSDR1* is involved in steroid synthesis and/or degradation in normal and neoplastic prostate epithelium, and, as such, it may be a key enzyme involved in maintaining intracellular balance of steroid hormones in these cells. Additional studies including the expression of *PSDR1* protein and the analysis of substrate specificity and kinetics are needed to address this question.

ACKNOWLEDGMENTS

We thank Steve Lasky, John Hall, and the Molecular Biotechnology Sequencing Facility for DNA sequencing support.

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